

Europäisches Patentamt
European Patent Office
Office européen des brevets



(11) **EP 0 811 637 B1**

(12)

EUROPEAN PATENT SPECIFICATION

(45) Date of publication and mention
of the grant of the patent:
23.08.2000 Bulletin 2000/34

(51) Int Cl.7: **C08F 10/00, C08F 2/34,
C08F 4/602, C08F 4/44**

(21) Application number: **97303810.2**

(22) Date of filing: **04.06.1997**

(54) Method for feeding a liquid catalyst to a fluidized bed polymerization reactor

Verfahren zur Zugabe eines Katalysators in flüssiger Form zu einem
Wirbelschichtpolymerisationsreaktor

Procédé d'alimentation d'un catalyseur liquide à un réacteur de polymérisation à lit fluidisé

(84) Designated Contracting States:
AT BE DE ES FR GB GR IT NL PT SE

• **Williams, Clark Curtis**
Charleston, West Virginia 25314 (US)

(30) Priority: **06.06.1996 US 659764**

(74) Representative: **Allard, Susan Joyce et al**
BOULT WADE TENNANT,
Verulam Gardens
70 Gray's Inn Road
London WC1X 8BT (GB)

(43) Date of publication of application:
10.12.1997 Bulletin 1997/50

(73) Proprietor: **UNION CARBIDE CHEMICALS &
PLASTICS TECHNOLOGY CORPORATION**
Danbury, Connecticut 06817-0001 (US)

(56) References cited:
EP-A- 0 584 609 EP-A- 0 593 083
EP-A- 0 659 773 EP-A- 0 764 665
US-A- 4 994 534 US-A- 5 453 471

(72) Inventors:
• **Goode, Mark Gregory**
Hurricane, West Virginia 25526 (US)

EP 0 811 637 B1

Note: Within nine months from the publication of the mention of the grant of the European patent, any person may give notice to the European Patent Office of opposition to the European patent granted. Notice of opposition shall be filed in a written reasoned statement. It shall not be deemed to have been filed until the opposition fee has been paid. (Art. 99(1) European Patent Convention).

Description**Field of the Invention**

5 [0001] This invention pertains to gas phase polymerization reaction in the presence of a liquid catalyst. More particularly, the present invention is directed to a method of feeding a liquid catalyst to a reactor so as to produce a particle lean zone in the reactor.

Background of the Invention

10 [0002] Liquid catalysts offer many advantages over conventional solid-supported catalysts. Unsupported liquid catalysts require less equipment and raw materials to make them and impart fewer impurities to the final polymer product. The activity of a liquid catalyst is not adversely influenced by the surface area of a support material. Additional advantages are encountered when a high-activity unsupported metallocene catalyst is used for polymerizations in a fluidized bed reactor.

15 [0003] The use of liquid or unsupported catalysts are disclosed in U.S. Patent No. 5,317,036. This patent is directed to the use of unsupported, soluble olefin polymerization catalysts, particularly metallocenes in liquid form, in gas phase reactions. These catalysts have a droplet size in the range of about 1 to about 1,000 μm .

20 [0004] However, when a liquid catalyst is employed in gas phase polymerization, several phenomena can occur. First, the soluble or liquid catalyst tends to deposition on the resin or polymer forming the fluidized bed which in turn leads to accelerated polymerization on the surface of the particles of the bed. As the coated resin particles increase in size, they are exposed to a higher fraction of catalyst solution or spray because of their increased cross-sectional dimensions. If too much catalyst is deposited on the polymer particles, they can grow so large that they cannot be fluidized thereby causing the reactor to be shut down.

25 [0005] Second, using liquid catalyst under conditions of high catalyst activity, e.g., a liquid metallocene catalyst, the initial polymerization rate is often so high that the newly formed polymer or resin particles can soften or melt, adhering to larger particles in the fluidized bed. This needs to be avoided or minimized to avert reactor shutdown.

[0006] On the other hand, if the polymer particles size is too small, entrainment can occur resulting in fouling of the recycle line, compressor, and cooler and increased static electricity leading to sheeting can result.

30 [0007] Accordingly, there is a need to control growth in polymer particle size when using a liquid catalyst in a gas phase fluidized polymerization process.

SUMMARY OF THE INVENTION

35 [0008] Surprisingly, the present invention provides a method for controlling growth in polymer particle sized when using a liquid catalyst in gas phase fluidized polymerization. The present invention provides a process for producing a polymer in a gas fluidized bed reaction vessel having a polymerization zone containing a bed of growing polymer particles under polymerization conditions which comprises:

- 40 (i) continuously or intermittently introducing one or more monomers, optionally at least one diene, optionally at least one inert particulate material, and optionally one or more inert gases into the polymerization zone;
- (ii) continuously or intermittently introducing into the reaction vessel an unsupported polymerization catalyst in liquid form without immediately contacting the bed of growing polymer particles so as to provide a particle lean zone;
- 45 (iii) continuously or intermittently withdrawing polymeric product from the reaction vessel; and
- (iv) continuously withdrawing unreacted monomers and gases from said polymerization and optionally compressing and cooling said gases while maintaining the temperature within the polymerization zone below the dew point of at least one monomer present in the polymerization zone.

BRIEF DESCRIPTION OF THE DRAWINGS

50 [0009] Figures 1-4 depict injection tube and nozzle configurations employed in the examples, which tubes and nozzles used to spray the liquid catalyst into a particle lean zone in the fluidized bed of the reactor.

DETAILED DESCRIPTION OF THE INVENTION

55 [0010] **Polymers.** Illustrative of the polymers which can be produced in accordance with the invention are the following: ethylene homopolymers and ethylene copolymers employing one or more $\text{C}_3\text{-C}_{12}$ alpha olefins; propylene homopolymers and propylene copolymers employing one or more $\text{C}_4\text{-C}_{12}$ alpha olefins; polyisoprene; polystyrene;

polybutadiene; polymers of butadiene copolymerized with styrene; polymers of butadiene copolymerized with acrylonitrile; polymers of isobutylene copolymerized with isoprene ethylene propylene rubbers and ethylene propylene diene rubbers; and polychloroprene.

[0011] Polymerization. The present invention is not limited to any specific type of gas phase polymerization reaction and can be carried out in a stirred or fluidized bed reactor. The invention can be carried out in a single reactor or multiple reactors (two or more reactors in series). In addition to well known conventional gas phase polymerizations processes, "condensed mode", including the so-called "induced condensed mode", and "liquid monomer" operation of a gas phase polymerization can be employed.

[0012] A conventional fluidized bed process for producing resins is practiced by passing a gaseous stream containing one or more monomers continuously through a fluidized bed reactor under reactive conditions in the presence of a polymerization catalyst. Product is withdrawn from the reactor. A gaseous stream of unreacted monomer is withdrawn from the reactor continuously and recycled into the reactor along with make-up monomer added to the recycle stream.

[0013] Condensed mode polymerizations are disclosed in U.S. Patent Nos. 4,543,399; 4,588,790; 5,352,749; and 5,462,999. Condensing mode processes are employed to achieve higher cooling capacities and, hence, higher reactor productivity. In these polymerizations a recycle stream, or a portion thereof, can be cooled to a temperature below the dew point in a fluidized bed polymerization process, resulting in condensing all or a portion of the recycle stream. The recycle stream is returned to the reactor. The dew point of the recycle stream can be increased by increasing the operating pressure of the reaction/recycle system and/or increasing the percentage of condensable fluids and decreasing the percentage of non-condensable gases in the recycle stream. The condensable fluid may be inert to the catalyst, reactants and the polymer product produced; it may also include monomers and comonomers. The condensing fluid can be introduced into the reaction/recycle system at any point in the system. Condensable fluids include saturated or unsaturated hydrocarbons. In addition condensable fluids of the polymerization process itself other condensable fluids, inert to the polymerization can be introduced to "induce" condensing mode operation. Examples of suitable condensable fluids may be selected from liquid saturated hydrocarbons containing 2 to 8 carbon atoms (e.g., propane, n-butane, isobutane, n-pentane, isopentane, neopentane, n-hexane, isohexane, and other saturated C₆ hydrocarbons, n-heptane, n-octane and other saturated C₇ and C₈ hydrocarbons, and mixtures thereof). Condensable fluids may also include polymerizable condensable comonomers such as olefins, alpha-olefins, diolefins, diolefins containing at least one alpha olefin, and mixtures thereof. In condensing mode, it is desirable that the liquid entering the fluidized bed be dispersed and vaporized quickly.

[0014] Liquid monomer polymerization mode is disclosed, in U.S. Patent No. 5,453,471, U.S. Serial No. 510,375, PCT 95/09826 (US) and PCT 95/09827 (US). When operating in the liquid monomer mode, liquid can be present throughout the entire polymer bed provided that the liquid monomer present in the bed is adsorbed on or absorbed in solid particulate matter present in the bed, such as polymer being produced or fluidization aids (e.g., carbon black) present in the bed, so long as there is no substantial amount of free liquid monomer present more than a short distance above the point of entry into the polymerization zone. Liquid mode makes it possible to produce polymers in a gas phase reactor using monomers having condensation temperatures much higher than the temperatures at which conventional polyolefins are produced. In general, liquid monomer process are conducted in a stirred bed or gas fluidized bed reaction vessel having a polymerization zone containing a bed of growing polymer particles. The process comprises continuously introducing a stream of one or more monomers and optionally one or more inert gases or liquids into the polymerization zone; continuously or intermittently introducing a polymerization catalyst into the polymerization zone; continuously or intermittently withdrawing polymer product from the polymerization zone; and continuously withdrawing unreacted gases from the zone; compressing and cooling the gases while maintaining the temperature within the zone below the dew point of at least one monomer present in the zone. If there is only one monomer present in the gas-liquid stream, there is also present at least one inert gas. Typically, the temperature within the zone and the velocity of gases passing through the zone are such that essentially no liquid is present in the polymerization zone that is not adsorbed on or absorbed in solid particulate matter.

[0015] Monomers. Monomers that can be employed in the process can include one or more C₂ to C₁₂ alpha-olefins; dienes such as those taught in U.S. Patent No. 5,317,036 to Brady et al. such as hexadiene, dicyclopentadiene, norbornadiene, and ethylidene norbornene; readily condensable monomers such as those taught in U.S. Patent No. 5,453,471 including isoprene, styrene, butadiene, isobutylene, and chloroprene and acrylonitrile.

[0016] Inert Particulate Materials. The process of the invention can optionally employ inert particulate materials as fluidization aids. These inert particulate materials can include carbon black, silica, talc, and clays, as well as inert polymeric materials. Carbon black has a primary particle size of 10 to 100 nanometers, an average size of aggregate of 0.1 to 10 µm, and a specific surface area of 30 to 1,500 m²/g. Silica has a primary particle size of 5 to 50 nanometers, an average size of aggregate of 0.1 to 10 µm, and a specific surface area of about 50 to 500 m²/g. Clay, talc, and polymeric materials have an average particle size of 0.01 to 10 µm and a specific surface area of 3 to 30 m²/g. These inert particulate materials are employed in amounts ranging from 0.3 to 80%, preferably 5 to 50%, based on the weight of the final product. They are especially useful for the polymerization of sticky polymers as disclosed in U.S. Patent

Nos. 4,994,534 and 5,304,588.

[0017] Chain Transfer Agents and Other Additives. Chain transfer agents, promoters, scavenging agents and other additives can be, and often are, employed in the polymerization process of the invention. Chain transfer agents are often used to control polymer molecular weight. Examples of these compounds are hydrogen and metal alkyls of the general formula M^3R^5g , where M^3 is a Group IA, IIA or IIIA metal, R^5 is an alkyl or aryl, and g is 1, 2, or 3. Preferably, a zinc alkyl is employed; and, of these, diethyl zinc is most preferred. Typical promoters include halogenated hydrocarbons such as $CHCl_3$, $CFCI_3$, CH_3CCl_3 , CF_2ClCCl_3 , and ethyltrichloroacetate. Such promoters are well known to those skilled in the art and are disclosed in, for example, U.S. Patent No. 4,988,783. Other organometallic compounds such as scavenging agents for poisons may also be employed to increase catalyst activity. Examples of these compounds include metal alkyls, such as aluminum alkyls, most preferably triisobutylaluminum. Some compounds may be used to neutralize static in the fluidized-bed reactor, others known as drivers rather than antistatic agents, may consistently force the static to from positive to negative or from negative to positive. The use of these additives is well within the skill of those skilled in the art. These additives may be added to the reaction zone separately or independently from the liquid catalyst if they are solids, or as part of the catalyst provided they do not interfere with the desired atomization. To be part of the catalyst solution, the additives should be liquids or capable of being dissolved in the catalyst solution.

[0018] Catalyst. Any type of polymerization catalyst may be used in the present process, provided it is stable and sprayable or atomizable when in liquid form. A single liquid catalyst may be used, or a liquid mixture of catalysts may be employed if desired. These catalysts are used with cocatalysts and promoters well known in the art. Examples of suitable catalysts include:

A. Ziegler-Natta catalysts, including titanium based catalysts such as those described in U.S. Patent Nos. 4,376,062 and 4,379,758. Ziegler-Natta catalysts are well known in the art, and typically are magnesium/titanium/electron donor complexes used in conjunction with an organoaluminum cocatalyst.

B. Chromium based catalysts such as those described in U.S. Patent Nos. 3,709,853; 3,709,954; and 4,077,904.

C. Vanadium based catalysts such as vanadium oxychloride and vanadium acetylacetonate, such as described in U.S. Patent No. 5,317,036.

D. Metallocene catalysts.

E. Cationic forms of metal halides, such as aluminum trihalides.

F. Cobalt catalysts and mixtures thereof such as those described in U.S. Patent Nos. 4,472,559 and 4,182,814.

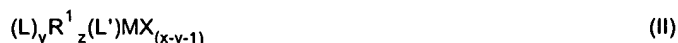
G. Nickel catalysts and mixtures thereof such as those described in U.S. Patent Nos. 4,155,880 and 4,102,817.

H. Rare Earth metal catalysts, i.e., those containing a metal having an atomic number in the Periodic Table of 57 to 103, such as compounds of cerium, lanthanum, praseodymium, gadolinium and neodymium. Especially useful are carboxylates, alcoholates, acetylacetonates, halides (including ether and alcohol complexes of neodymium trichloride), and allyl derivatives of such metals. Neodymium compounds, particularly neodymium neodecanoate, octanoate, and versatate, are the most preferred rare earth metal catalysts. Rare earth catalysts are used to produce polymers polymerized using butadiene or isoprene.

[0019] Preferred among these different catalyst systems are catalyst compositions comprising a metallocene catalyst in liquid form and an activating cocatalyst. The practice of this invention is not limited to any particular class or kind of metallocene catalyst. Accordingly, the catalyst composition may comprise any unsupported metallocene catalyst useful in slurry, solution, bulk, or gas phase olefin polymerization. One or more than one metallocene catalyst may be employed. For example, as described in U.S. Patent No. 4,530,914, at least two metallocene catalysts may be used in a single catalyst composition to achieve a broadened molecular weight distribution polymer product.

[0020] Metallocene catalysts are organometallic coordination complexes of one or more π -bonded moieties in association with a metal atom from Groups IIIB to VIII or the rare earth metals of the Periodic Table.

[0021] Bridged and unbridged mono-, bis-, and tris-cycloalkadienyl/metal compounds are the most common metallocene catalysts, and generally are of the formula:

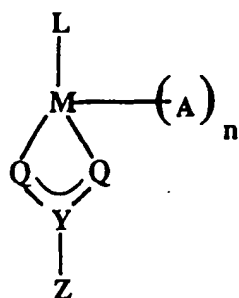


wherein M is a metal from groups IIIB to VIII of the Periodic Table; L and L' are the same or different and are π -bonded ligands coordinated to M, preferably cycloalkadienyl groups such as cyclopentadienyl, indenyl, or fluorenyl groups optionally substituted with one or more hydrocarbyl groups containing 1 to 20 carbon atoms; R^1 is a C_1 - C_4 substituted or unsubstituted alkylene radical, a dialkyl or diaryl germanium or silicon, or an alkyl or aryl phosphine or amine radical bridging L and L'; each X is independently hydrogen, an aryl, alkyl, alkenyl, alkylaryl, or arylalkyl radical having 1-20

carbon atoms, a hydrocarboxy radical having 1-20 carbon atoms, a halogen, $R^2CO_2^-$, or $R^2_2NCO_2^-$, wherein each R^2 is a hydrocarbyl group containing 1 to about 20 carbon atoms; n and m are each 0, 1, 2, 3, or 4; y is 0, 1, or 2; x is 1, 2, 3, or 4 depending upon the valence state of M; z is 0 or 1 and is 0 when y is 0; and $x-y \geq 1$.

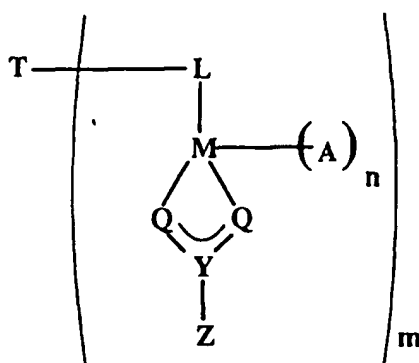
[0022] Illustrative examples of metallocene catalysts represented by formula II are dialkyl metallocenes such as bis(cyclopentadienyl)titanium dimethyl, bis(cyclopentadienyl)titanium diphenyl, bis(cyclopentadienyl)zirconium dimethyl, bis(cyclopentadienyl)zirconium diphenyl, bis(cyclopentadienyl)hafnium methyl and diphenyl, bis(cyclopentadienyl)titanium di-neopentyl, bis(cyclopentadienyl)zirconium di-neopentyl, bis(cyclopentadienyl)titanium dibenzyl, bis(cyclopentadienyl)zirconium dibenzyl, bis(cyclopentadienyl)vanadium dimethyl; mono alkyl metallocenes such as bis(cyclopentadienyl)titanium methyl chloride, bis(cyclopentadienyl)titanium ethyl chloride, bis(cyclopentadienyl)titanium phenyl chloride, bis(cyclopentadienyl)zirconium methyl chloride, bis(cyclopentadienyl)zirconium ethyl chloride, bis(cyclopentadienyl)zirconium phenyl chloride, bis(cyclopentadienyl)titanium methyl bromide; trialkyl metallocenes such as cyclopentadienyl titanium trimethyl, cyclopentadienyl zirconium triphenyl, and cyclopentadienyl zirconium trineopentyl, cyclopentadienyl zirconium trimethyl, cyclopentadienyl hafnium triphenyl, cyclopentadienyl hafnium trineopentyl, and cyclopentadienyl hafnium trimethyl; monocyclopentadienyl titanocenes such as, pentamethylcyclopentadienyl titanium trichloride, pentaethylcyclopentadienyl titanium trichloride; bis(pentamethylcyclopentadienyl) titanium diphenyl, the carbene represented by the formula bis(cyclopentadienyl)titanium= CH_2 and derivatives of this reagent; substituted bis(cyclopentadienyl)titanium (IV) compounds such as: bis(indenyl)titanium diphenyl or dichloride, bis(methylcyclopentadienyl)titanium diphenyl or dihalide; dialkyl, trialkyl, tetraalkyl and pentaalkyl cyclopentadienyl titanium compounds such as bis(1,2-dimethylcyclopentadienyl)titanium diphenyl or dichloride, bis(1,2-diethylcyclopentadienyl)titanium diphenyl or dichloride; silicon, phosphine, amine or carbon bridged cyclopentadiene complexes, such as dimethyl silyldicyclopentadienyl titanium diphenyl or dichloride, methyl phosphine dicyclopentadienyl titanium diphenyl or dichloride, methylenedicyclopentadienyl titanium diphenyl or dichloride and other dihalide complexes, and the like; as well as bridged metallocene compounds such as isopropyl(cyclopentadienyl)(fluorenyl)zirconium dichloride, isopropyl(cyclopentadienyl)(octahydrofluorenyl)zirconium dichloride, diphenylmethylene(cyclopentadienyl)(fluorenyl)zirconium dichloride, diisopropylmethylene(cyclopentadienyl)(fluorenyl)-zirconium dichloride, diisobutylmethylene(cyclopentadienyl)(fluorenyl)zirconium dichloride, ditertbutylmethylene(cyclopentadienyl)(fluorenyl)zirconium dichloride, cyclohexylidene(cyclopentadienyl)(fluorenyl)zirconium dichloride, diisopropylmethylene(2,5-dimethylcyclopentadienyl)(fluorenyl)zirconium dichloride, isopropyl(cyclopentadienyl)(fluorenyl)hafnium dichloride, diphenylmethylene(cyclopentadienyl)(fluorenyl)hafnium dichloride, diisopropylmethylene(cyclopentadienyl)(fluorenyl)hafnium dichloride, ditertbutylmethylene(cyclopentadienyl)(fluorenyl)hafnium dichloride, cyclohexylidene(cyclopentadienyl)(fluorenyl)hafnium dichloride, diisopropylmethylene(2,5-dimethylcyclopentadienyl)(fluorenyl)hafnium dichloride, isopropyl(cyclopentadienyl)(fluorenyl)titanium dichloride, diphenylmethylene(cyclopentadienyl)(fluorenyl)titanium dichloride, diisopropylmethylene(cyclopentadienyl)(fluorenyl)titanium dichloride, diisobutylmethylene(cyclopentadienyl)(fluorenyl)titanium dichloride, ditertbutylmethylene(cyclopentadienyl)(fluorenyl)titanium dichloride, cyclohexylidene(cyclopentadienyl)(fluorenyl)titanium dichloride, diisopropylmethylene(2,5 dimethylcyclopentadienyl fluorenyl)titanium dichloride, racemic-ethylene bis (1-indenyl) zirconium (IV) dichloride, racemic-ethylene bis (4,5,6,7-tetrahydro-1-indenyl) zirconium (IV) dichloride, racemic-dimethylsilyl bis (1-indenyl) zirconium (IV) dichloride, racemic-dimethylsilyl bis (4,5,6,7-tetrahydro-1-indenyl) zirconium (IV) dichloride, racemic-1,1,2,2-tetramethylsilanylene bis (1-indenyl) zirconium (IV) dichloride, racemic-1,1,2,2-tetramethylsilanylene bis (4,5,6,7-tetrahydro-1-indenyl) zirconium (IV) dichloride, ethylidene (1-indenyl tetramethylcyclopentadienyl) zirconium (IV) dichloride, racemic-dimethylsilyl bis (2-methyl-4-t-butyl-1-cyclopentadienyl) zirconium (IV) dichloride, racemic-ethylene bis (1-indenyl) hafnium (IV) dichloride, racemic-ethylene bis (4,5,6,7-tetrahydro-1-indenyl) hafnium (IV) dichloride, racemic-dimethylsilyl bis (1-indenyl) hafnium (IV) dichloride, racemic-dimethylsilyl bis (4,5,6,7-tetrahydro-1-indenyl) hafnium (IV) dichloride, racemic-1,1,2,2-tetramethylsilanylene bis (1-indenyl) hafnium (IV) dichloride, racemic-1,1,2,2-tetramethylsilanylene bis (4,5,6,7-tetrahydro-1-indenyl) hafnium (IV), dichloride, ethylidene (1-indenyl-2,3,4,5-tetramethyl-1-cyclopentadienyl) hafnium (IV) dichloride, racemic-ethylene bis (1-indenyl) titanium (IV) dichloride, racemic-ethylene bis (4,5,6,7-tetrahydro-1-indenyl) titanium (IV) dichloride, racemic-dimethylsilyl bis (1-indenyl) titanium (IV) dichloride, racemic-dimethylsilyl bis (4,5,6,7-tetrahydro-1-indenyl) titanium (IV) dichloride, racemic-1,1,2,2-tetramethylsilanylene bis (1-indenyl) titanium (IV) dichloride, racemic-1,1,2,2-tetramethylsilanylene bis (4,5,6,7-tetrahydro-1-indenyl) titanium (IV) dichloride, and ethylidene (1-indenyl-2,3,4,5-tetramethyl-1-cyclopentadienyl) titanium (IV) dichloride.

[0023] Particularly preferred metallocene catalysts have one of the following formulas (III or IV):



(III)

or



(IV)

wherein:

M is a metal from groups IIIB to VIII, preferably Zr or Hf;

L is a substituted or unsubstituted, π -bonded ligand coordinated to M, preferably a substituted cycloalkadienyl ligand;

each Q is independently selected from the group consisting of -O-, -NR³-, -CR³₂- and -S-, preferably oxygen;

Y is either C or S, preferably carbon;

Z is selected from the group consisting of -OR³, -NR³₂, -CR³₃, -SR³, -SiR³₃, -PR³₂, and -H, with the proviso that when Q is -NR³- then Z is selected from the group consisting of -OR³, -NR³₂, -SR³, -SiR³₃, -PR³₂, and -H, preferably

Z is selected from the group consisting of -OR³, -CR³₃, and -NR³₂;

n is 1 or 2;

A is a univalent anionic group when n is 2 or A is a divalent anionic group when n is 1, preferably A is a carbamate, carboxylate or other heteroallyl moiety described by Q, Y and Z combination; and

each R³ is independently a group containing carbon, silicon, nitrogen, oxygen, and/or phosphorus and one or more R³ groups may be attached to the L substituent, preferably R³ is a hydrocarbon group containing from 1 to 20 carbon atoms, most preferably an alkyl, cycloalkyl or an aryl group;

T is a bridging group selected from the group consisting of alkylene or arylene groups containing from 1 to 10 carbon atoms optionally substituted with carbon or heteroatoms, germanium, silicon and alkyl phosphine; and

m is 1 to 7, preferably 2 to 6, most preferably 2 or 3.

[0024] The supportive substituent formed by Q, Y and Z is a uncharged polydentate ligand exerting electronic effects due to its high polarizability, similar to the cyclopentadienyl group. In the most preferred embodiments of this invention, the disubstituted carbamates,



15



25

30

35



45

50

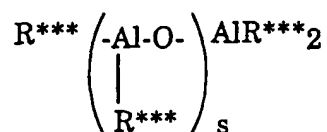
55

7

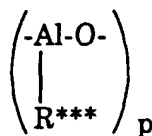
Z'	Cp	Y'	X'	M
dimethylsilyl methylphenylsilyl diphenylsilyl tetramethylethylene ethylene diphenylmethylene	cyclopentadienyl fluorenyl indenyl tetramethylcyclopentadienyl	t-butylamido phenylamido cyclohexylamido oxo	chloride methyl	titanium zirconium hafnium

[0029] The activating cocatalyst is capable of activating the metallocene catalyst. Preferably, the activating cocatalyst is one of the following: (a) branched or cyclic oligomeric poly(hydrocarbyl-aluminum oxide)s which contain repeating units of the general formula $-(Al(R^*)O)-$, where R^* is hydrogen, an alkyl radical containing from 1 to about 12 carbon atoms, or an aryl radical such as a substituted or unsubstituted phenyl or naphthyl group; (b) ionic salts of the general formula $[A^+][BR^{**4-}]$, where A^+ is a cationic Lewis or Bronsted acid capable of abstracting an alkyl, halogen, or hydrogen from the metallocene catalysts, B is boron, and R^{**} is a substituted aromatic hydrocarbon, preferably a perfluorophenyl radical; and (c) boron alkyls of the general formula BR^{**3} , where R^{**} is as defined above.

[0030] Preferably, the activating cocatalyst is an aluminoxane such as methylaluminoxane (MAO) or modified methylaluminoxane (MMAO), or a boron alkyl. Aluminoxanes are preferred and their method of preparation is well known in the art. Aluminoxanes may be in the form of oligomeric linear alkyl aluminoxanes represented by the formula:



or oligomeric cyclic alkyl aluminoxanes of the formula:



wherein s is 1-40, preferably 10-20; p is 3-40, preferably 3-20; and R^{***} is an alkyl group containing 1 to 12 carbon atoms, preferably methyl or an aryl radical such as a substituted or unsubstituted phenyl or naphthyl radical. In the case of MAO, R^{***} is methyl, whereas in MMAO, R^{***} is a mixture of methyl and C2 to C12 alkyl groups wherein methyl comprises about 20 to 80 percent by weight of the R^{***} group.

[0031] The amount of activating cocatalyst and metallocene catalyst usefully employed in preparation of the catalyst composition, whether the catalyst composition is formed in situ as it is being introduced into the reaction zone or formed prior to introduction into the reaction zone, can vary over a wide range. When the cocatalyst is a branched or cyclic oligomeric poly(hydrocarbylaluminum oxide), the mole ratio of aluminum atoms contained in the poly(hydrocarbylaluminum oxide) to metal atoms contained in the metallocene catalyst is generally in the range of from 2:1 to 100,000:1, preferably in the range of from 10:1 to 10,000:1, and most preferably in the range of from 50:1 to 2,000:1. When the cocatalyst is an ionic salt of the formula $[A^+][BR^{*4-}]$ or a boron alkyl of the formula BR^{*3} , the mole ratio of boron atoms contained in the ionic salt or the boron alkyl to metal atoms contained in the metallocene catalyst is generally in the range of from 0.5:1 to 10:1, preferably in the range of from 1:1 to 5:1.

[0032] The liquid catalyst can be composed of one or more of metal compounds in combination with one or more co-catalysts. Alternatively, all or a portion of the co-catalyst can be fed separately from the metal compound(s) to the reactor. Promoters associated with any particularly polymerization are usually added to the reactor separately from the co-catalyst and/or metal compound(s).

[0033] If the metal compound and/or the co-catalyst occurs naturally in liquid form, it can be introduced "neat" into

the particle lean zone. More likely, the liquid catalyst is introduced into the particle lean zone as a solution (single phase, or "true solution" using a solvent to dissolve the metal compound and/or co-catalyst), an emulsion (partially dissolving the catalyst components in a solvent), suspension, dispersion, or slurry (each having at least two phases). Preferably, the liquid catalyst employed is a solution or an emulsion, most preferably a solution. As used herein, "liquid catalyst" or "liquid form" includes neat, solution, emulsion, and dispersions of the transition metal or rare earth metal component(s) of the catalyst and/or co-catalyst.

[0034] The solvents that which can be utilized to form solutions of the soluble, unsupported transition metal and/or rare earth metal polymerization catalyst compounds are inert solvents, preferably non-functional hydrocarbon solvents, and may include aliphatic hydrocarbons such as butane, isobutane, ethane, propane, pentane, isopentane, hexane, heptane, octane, decane, dodecane, hexadecane and octadecane; alicyclic hydrocarbons such as cyclopentane, methylcyclopentane, cyclohexane, cyclooctane, norbornane and ethylcyclohexane; aromatic hydrocarbons such as benzene, toluene, ethylbenzene, propylbenzene, butylbenzene, xylene and tetrahydrofuran; and petroleum fractions such as gasoline, kerosene and light oils. Likewise, halogenated hydrocarbons such as methylene chloride and chlorobenzene may also be utilized. By "inert" is meant that the material being referred to is non-deactivating in the polymerization reaction zone under the conditions of gas phase polymerization and is non-deactivating with the catalyst in or out of the reaction zone. By "non-functional", it is meant that the solvents do not contain groups such as strong polar groups which can deactivate the active catalyst metal sites.

[0035] The concentration of the catalyst and/or co-catalyst that is in solution that is provided to the lean particle zone may be as high as the saturation point of the particular solvent being used. Preferably, the concentration is in the range of from 0.01 to 10,000 millimoles/liter. Of course, if the catalyst and/or co-catalyst is being used in its neat form, i.e., in its liquid state with no solvent, it will be comprised of essentially pure catalyst and/or co-catalyst, respectively.

[0036] The size of the droplets formed when introducing the catalyst system into the reactor is generally determined by the manner and place in which the catalyst is introduced. It is desirable to use a means of introduction which is able to provide liquid droplets in the particle lean zone having an average diameter which is in the range of from 0.1 to 1000 μm , preferably within a range of 0.1 to 500 μm , most preferably ranging from 1 to 150 μm . A narrow distribution of droplet size in a lower or mid range of 10 to 100 can prevent the formation of large agglomerates resulting from large droplets and the formation of fines resulting from small droplets. Under many conditions, however, a wide droplet size distribution is acceptable as the smaller droplets can agglomerate to some degree with the resin in the reactor and large droplets can form larger particles of up to 0.25 which can be readily fluidized as long as the particle fraction is low enough, preferably less than 10 % and more preferably less than 2% by weight of the total resin in the bed.

[0037] Establishing a Resin Particle Lean Zone. A resin particle lean zone can be established in the reactor by feeding the catalyst in any manner such that the catalyst droplets do not immediately contact a substantial portion of the resin particles of the fluidized bed. The droplets of the unsupported catalyst in liquid form are introduced without immediately contacting growing polymer particles of the bed so as to provide an average polymer particle size (APS) ranging from 0.25 to 1.52 mm (0.01 to 0.06 inches). Generally, the particle density in the particle lean zone is at least 10 times lower than that in the fluidized bed. As disclosed in U.S. Patent No. 5,317,036, a liquid, unsupported catalyst is typically dispersed in a solvent such as isopentane and introduced into the fluidized bed using an inert carrier gas such as nitrogen. In the time period elapsing when the liquid catalyst in droplet form leaves the nozzle and contacts the particles in the bed, new polymer particles are formed. In the present invention, the time between the droplet leaving the nozzle and its contacting the particles in the bed ranges from 0.01 seconds to 60 seconds, preferably 0.01 to 30 seconds, and, most preferably, is 0.01 seconds to 5 seconds.

[0038] A particle lean zone may be a section of the reactor which normally does not contain the fluidized bed, such as the disengaging section, the gas recirculation system, or the area below the distributor plate. The particle lean zone may also be created by deflecting resin away from the catalyst spray with a stream of gas.

[0039] In a preferred embodiment of the present invention, the liquid catalyst in a carrier gas (e.g., nitrogen, argon, alkane, or mixtures thereof) is surrounded by at least one gas which serves to move or deflect resin particles of the bed out of the path of the liquid catalyst as it enters the fluidization zone and away from the area of catalyst entry, thereby providing a particle lean zone. In a particularly preferred embodiment, the liquid catalyst in the carrier gas is surrounded by at least two gases, the first gas serving primarily to deflect resin particles of the bed out of the path of the liquid catalyst and the second gas primarily prevents the injection tube or nozzle tip from getting clogged. As used in this application, when the liquid catalyst in the carrier gas is surrounded by two gases, the catalyst is considered to be shrouded. The first or particle-deflecting gas and the second or tip-cleaning gas can each be selected from the group consisting of recycle gas, monomer gas, chain transfer gas (e.g., hydrogen), insert gas or mixtures thereof. Preferably the particle-deflecting gas is all or a portion of the recycle gas and the tip-cleaning gas is all or a portion of a monomer (e.g., ethylene or propylene) employed in the process.

[0040] Liquid catalyst in a carrier gas, particle-deflecting gas, and, when employed, the tip-cleaning gas can be introduced into the reactor at the same velocities to establish a particle lean zone. However, it is preferred that they enter the fluidization zone at differing velocities. Preferably, the liquid catalyst in the carrier gas is introduced at a

velocity of from 15 to 120 m/s (50 ft/s to 400 ft/s); the particle-deflecting gas is introduced at a velocity of from 3 to 46 m/s (10 ft/s to 150 ft/s) and, when employed, the tip-cleaning gas ranges in velocity from 15 to 76 m/s (50 ft/s to 250 ft/s). Preferably, the pressure of the particle-deflecting gas, and, when employed, the tip-cleaning gas is 70 to 340 kPa gauge (10 to 50 psig), preferably 140 to 210 kPa gauge (20 to 30 psig), higher than the pressure of the gas in the polymerization zone of the reactor. Typically, the particle-deflecting gas pressure ranges from 70 to 340 kPa gauge (10 to 50 psig); the tip-cleaning gas pressure, when employed, ranges from 340 to 1700 kPa gauge (50 to 250 psig); and the liquid catalyst/carrier gas pressure ranges from 340 to 1700 kPa gauge (50 to 250 psig). When the particle-deflecting gas is the recycle gas, it is a portion comprising 5 to 25 percent of the total recycle flow and is preferably removed from the discharge side of the compressor. When the tip-cleaning gas is the monomer gas, it is a portion comprising 2 to 40 percent of the total monomer flow. The particle-deflecting gas and the tip-cleaning gas can also optionally contain one or more antifoulants or antistatic agents known to those skilled in the art. While inert gases can be employed in the present invention as the particle-deflecting and tip-cleaning gases, they can be impractical because they require increased reactor venting, thereby decreasing efficiency of monomer usage and increasing cost.

[0041] Unsupported liquid catalyst can be introduced into the polymerization zone from the side, top, or bottom of the reactor. Side feeding the liquid catalyst is generally preferred, since it requires no or little modification of a conventional commercial reactor. When the liquid, unsupported catalyst is fed from a side location into the fluidization or polymerization zone of the reactor, it, along with the particle-deflecting gas and optional tip-cleaning gas, preferably enters the bed from a position that is 10 percent to 40 percent of the distance from the distributor plate to the top of the bed, most preferably 15 percent to 25 percent of the distance from the distributor plate to the top of the bed. When the liquid, unsupported catalyst is fed from the bottom of the reactor along with the particle-deflecting gas and optional tip-cleaning gas, it preferably enters the fluidized bed from a position that is at or near the center of the distributor plate in the bottom of the reactor to provide a particle lean zone. When the unsupported, liquid catalyst is introduced from a location in the top of the reactor, it is preferred that it enter in such a manner so as to avoid polymerization in the expanded zone of the reactor, and, therefore, is released in the reactor at the top or just immediately above the fluidized bed. This allows the catalyst droplets to additionally coat fines which can accumulate as dust above the top of the fluidized bed.

[0042] Any catalyst delivery system that is capable of atomizing the liquid catalyst into droplets of the desired size and distribution and avoids plugging of the tip or nozzle can be employed in the present invention. One embodiment of a catalyst delivery system comprises a particle-deflecting gas tube enclosing an optional tip-cleaning gas tube which in turn encloses a catalyst injection tube. The particle-deflecting gas tube has a sufficient inside diameter for the insertion or mounting of the tip-cleaning gas tube. For a commercial fluidized bed reactor, typically the particle-deflecting gas tube has an inside diameter ranging from 5 to 30 cm (2 inches to 12 inches), preferably 10 to 15 cm (4 to 6 inches). The optional tip-cleaning gas tube, has an outside diameter capable of fitting inside the particle-deflecting gas tube. For a conventional reactor, typically the tip-cleaning gas tube has an inside diameter ranging from 1.3 to 3.8 cm (0.5 inches to 1.5 inches), preferably 1.9 to 3.2 cm (0.75 to 1.25 inches).

[0043] The particle-deflecting gas tube can be flush with the inside wall of the reactor or lead edge (top surface) of the distributor plate, or, preferably, it can be extended beyond the inside wall of the reactor or lead edge of the distributor plate into the fluidization zone. Preferably the particle-deflecting gas tube is flush with the inside wall or top of the distributor plate. When employed the tip-cleaning gas tube can be positioned flush with, extended beyond, or recessed in the particle-deflecting gas tube. Preferably the tip-cleaning gas tube is flush with or recessed in the particle-deflecting gas tube. Most preferably the tip-cleaning gas tube is flush with the particle-deflecting gas tube.

[0044] The catalyst injection tube or nozzle can be housed within the particle-deflecting gas tube, but is preferably housed within the tip-cleaning gas tube which is inside the particle-deflecting gas tube. Preferably the catalyst injection tube or nozzle is tapered at its tip to a fine or knife edge to minimize surface area for injector fouling and convenient entry to the reactor vessel. The catalyst injection tube or nozzle is secured or anchored to the inner wall of the particle-deflecting gas tube or preferably to the tip-cleaning gas tube by means of one or more fins or flanges. Stainless steel injection tubing and pneumatic spray nozzles are commercially available in a wide range of internal diameters and thicknesses such that tubing or nozzle size can easily be matched to the amount of catalyst solution feed. For a commercial-size fluidized bed reactor, tubing and nozzles having about a 3 mm (1/8-inch) inside diameter are employed. The orifice diameter in the spray nozzle tip is in the range of from 0.3 to 6.4 mm (0.01 inch to 0.25 inch), preferably from 0.5 to 3.8 mm (0.02 inch to 0.15 inch). The orifice diameter of the tip of the injection tube is between 1.3 to 6.4 mm (0.05 inch to 0.25 inches), preferably between 2.5 to 5.1 mm (0.1 inch to 0.2 inches). Suitable nozzles can be obtained from Spraying Systems Corporation (Wheaton, IL) and can include the 1/8 JJ Series having standard and customized configurations. For a given liquid catalyst and reactor polymerization conditions the catalyst liquid feed rates can be adjusted by one skilled in the art to obtain the desired droplet size and distribution. The catalyst injection tube or nozzle can be located flush, extended, or recessed with respect to the leading tip edge of the particle-deflecting gas tube and/or optional tip-cleaning gas tube.

[0045] In the absence of the tip-cleaning gas tube, the catalyst injection tube or nozzle can be located flush, extended,

or recessed with respect to the leading tip edge of the particle-deflecting gas tube. Preferably the catalyst injection tube or nozzle is located flush or extended with respect to the leading tip edge of the particle-deflecting gas tube in the absence of the tip-cleaning gas tube. Most preferably it is located flush in the particle-deflecting gas tube. When a tip-cleaning gas tube is employed in conjunction with the particle-deflecting gas tube, the catalyst injection tube or nozzle is extended beyond the leading edge of the tip-cleaning gas tube or flush with the leading edge to the tip-cleaning gas tube. Preferably, the catalyst injection tube or nozzle is extended 5 to 10 cm (2 to 4 inches) beyond the leading edge of the tip-cleaning gas tube, but recessed with respect to the particle-deflecting gas tube.

[0046] All references cited herein are incorporated by reference.

[0047] Whereas the scope of the invention is set forth in the appended claims, the following specific examples illustrate certain aspects of the present invention. The examples are set forth for illustration only and are not to be construed as limitations on the invention, except as set forth in the claims. All parts and percentages are by weight unless otherwise specified.

EXAMPLES

Preparation of Liquid Metallocene Catalyst.

[0048] Indene (available from Nippon Steel) and $\text{Zr}(\text{NEt}_2)_4$ (available from Boulder Scientific) were reacted in stoichiometric amounts to form $\text{IndZr}(\text{NEt}_2)_3$. Gaseous carbon dioxide was bubbled through a 0.05 Molar solution of $\text{IndZr}(\text{NEt}_2)_3$ for fifteen minutes while being cooled with an ice bath. The solution was then sparged for fifteen minutes with nitrogen to remove excess carbon dioxide.

Polymerization in Pilot-scale Fluidized Bed Reactor Using Liquid Metallocene Catalyst.

[0049] In Examples 1 to 5, polymerizations were conducted in pilot-scale fluidized-bed reactors of 36 or 46 cm (14- or 18-inch) nominal diameter having a cylindrical section containing the fluidized-bed, an expanded section for disengagement of entrained resin, a gas compressor, a heat exchanger used for control of the resin temperature, and a product discharge system. The flowrate of reactant and inert gas through the reactor system was adjusted by a ball valve on the recycle line to provide a superficial gas velocity in the polymerization zone of 0.46 to 0.56 m/s (1.5 to 1.8 ft/s). The gas was circulated through the reactor at a rate between 3.6 and 5.9 tonne/hr (8,000 and 13,000 lbs/hr). Resin was produced at a rate between 14 and 18 Kg/hr (30 and 40 lbs/hr) to provide a resin residence time of 3 to 4 hours.

[0050] The liquid catalyst prepared above was metered to the reactor with a piston pump. Alumoxane cocatalyst was pressurized under 500-psi nitrogen and metered with a motor valve. The catalyst and cocatalyst were contacted in standard 3.2 or 6.4 mm (1/8 or 1/4") tubing tees and allowed to mix for several minutes before being diluted with isopentane. Mixing times varied between a few minutes and an hour, depending on the activation requirements of the catalyst. When an injection tube was used, the catalyst solution was mixed with nitrogen atomization gas just upstream of the reactor. When nozzles were used, the catalyst/cocatalyst mixture was delivered to the nozzle separately from the nitrogen. Catalyst concentrations and feed rates, and cocatalyst feed rates are listed in Table I of the examples. The alumoxane was supplied by Akzo-Nobel and is designated as MMAO type 3A. This was mixed with an equal weight of purified isopentane to provide a 4.4 wt % Al solution, for convenience in handling.

[0051] Reactor startup was accomplished by charging a seed bed and drying to less than 5 ppm water. The reactor was filled with monomer and inert gas to the desired levels and heated to the desired temperature. A nitrogen purge flow was maintained on the nozzles and/or injection tubes during this time. Cocatalyst flow was initiated, and once flowrate control was established, the catalyst feed was begun. Polymerization initiated immediately after the catalyst/cocatalyst mixture reached the reactor.

[0052] The average particle size was determined by a sieve analysis using ASTM D-1921 Method A.

[0053] Liquid catalyst was sprayed into the pilot-scale reactor with three different injectors: (1) a commercial nozzle adapted from reactor use, (2) the same nozzle with an elongated air cap, and (3) a tapered injection tube.

[0054] The commercial spray nozzle (1) consisted of a body, a liquid cap, and a tapered spraying tip. The liquid cap was purchased from Spraying Systems Co (1/8 JJ series #PFJ1650). Liquid catalyst along with nitrogen for atomization was fed through coaxial tubing to the body of the nozzle, which housed the liquid cap. The body was 96 mm (5/8 inches) in diameter and 28.58 mm (1.125 inches) long. The spray was formed at the liquid cap and fed through the air cap to the reactor. The air cap was either purchased from Spraying Systems (#PAJ67147) as depicted in Figure 1, or was machine in-house to an elongated tapered air cap (2) as depicted in Figure 2. The nozzle tip was 3.8 cm (1.5 inches) long and was tapered over the last 1.3 cm (0.5 inches) to reduce the accumulation of resin near the spray orifice. The orifice diameter was 1 mm (0.04 inches). The air cap was held onto the reactor with a retaining nut (Spraying Systems CP12582). The third type of injector, was a 3.2 mm (1/8 inch)-OD x 1.4 mm (0.055-inch)-ID stainless steel tube which

was tapered over the last 1.3 cm (1/2 inch) to produce a sharp edge. This design produced in-house is depicted in Figure 3.

[0055] In examples 2 and 4, the spray nozzle illustrated in Figure 2 was enclosed in a shroud to avoid polymerization near the nozzle orifice, as shown in Figure 4. The shroud was made from 1.9 cm (0.75 inch) OD stainless steel tubing and was capped at the end with a machined tapered tip. The tip-cleaning gas was fed from behind the spray nozzle and passed around the 1.6 cm (5/8 inch) adapter piece past the tip of the nozzle.

[0056] For Examples 1 and 5, a commercial Spraying Systems Co. 1/8 JJ nozzle was used with a J-1650 liquid cap and a PAJ67147 air cap. The nozzle body was modified in-house to allow coaxial introduction of catalyst. For Examples 2 and 4, a customized air cap nozzle, 1.5 inches long with a tapered tip was used. For these examples, the nozzles were located within a 1.9 cm (0.75-inch) tip-cleaning gas tube. For Example 3, a tapered 3.2 mm (1/8-inch) injection tube was used.

[0057] In Example 1 (Comparative) (using a commercial spray nozzle), catalyst was fed through a spray nozzle at the side of the reactor, one foot above the distributor plate without providing a particle lean zone. The average particle size increased from 0.4 to 1.3 mm (0.016 to 0.0511 inches) soon after the reaction began, indicating that most of the new particles were agglomerating with the particles in the reactor.

[0058] In Example 5 (employing a commercial spray nozzle), catalyst was sprayed at the top of the fluidized-bed into the expanded section to produce small particles.

[0059] In Example 4, catalyst was fed in a particle deflecting gas composed of recycle gas (193 kg/hr-426 lbs/hr) from a standpipe in the center of the distributor plate (Figure 4). A stable APS of 1 mm (0.04 inches) was obtained.

[0060] In Examples 2 (a tapered spray nozzle with a N₂ shroud) and 3 (an injection tube in a particle deflecting gas in the absence of a shroud), catalyst was fed surrounded by a particle deflecting gas of recycle gas into the side of the reactor about 30 cm (1 foot) above the plate in the absence of a tip-cleaning gas. This provided stable APS in the 0.64 to 0.71 mm (0.025 to 0.028 inch) range.

[0061] By feeding the catalyst in the stream of recycle gas, as done in Examples 2, 3, and 4, a particle-lean zone is provided in the jet or the resulting bubbles, so that the droplets can polymerize without adhering to the resin in the bed.

TABLE 1

Catalyst Feeding Location	Ex. 1	Ex. 2	Ex. 3	Ex. 4	Ex. 5
	Side of reactor without jet gas	Side of reactor with recycle gas jet	Side of reactor with recycle gas jet	Center of distributor plate with recycle gas jet	Top of fluidized bed
Catalyst Molarity	0.025	0.05	0.05	0.05	0.025
Catalyst Feedrate (cc/hr)	10	10	5	12	5
MMAO feedrate (cc/hr)	100	120	80	180	100
iCS feedrate kg/hr (lbs/hr)	0.68 (1.5)	0	0	0	0.68 (1.5)
N ₂ feedrate (lbs/hr)	3.2 (7)	3.4 (7.5)	3.2 (7)	1.8(4)	4.5 (10)
Temperature (°C)	80	80	80	80	80
Pressure MPa gauge (psig)	2.41 (350)	2.41 (350)	2.53 (367)	2.52 (365)	2.41 (350)
Ethylene pressure MPa (psia)	1.38 (200)	1.38 (200)	1.52 (220)	1.38 (200)	1.52 (220)
C ₆ /C ₂ (molar ratio)	0.01 to 0.012	0.012	0.026	0.02	0.003
Shroud N ₂ rate kg/hr (lbs/hr)*	NA	4.5 (10)	NA	3.6 (8)	NA
Recycle jet rate kg/hr (lbs/hr)†	NA	685.8 (1512)	725.7 (1600)	193 (426)	NA
Particle size	Increased from 0.41 to 1.30 mm (0.016 to 0.0511")	Stable at 0.71mm (0.028")	Stable at 0.64mm (0.025")	Stable at 1mm (0.04")	0.2mm (0.008") spherical particles recovered from disengaging section

* Used as a tip cleaning gas.

† Used as a particle deflecting gas.

Polymerizations in a Commercial-scale Reactor with Solution Metallocene Catalyst

[0062] The indenyl zirconium tris-diethyl carbamate catalyst was used to polymerize a ethylene-hexene copolymer (density 0.922, melt index 3) in a commercial-scale fluidized bed reactor. The reactor was nominally 2.4m (8 feet) in diameter and was operated with a bed height of 11.6 to 12.8m (38 to 42 feet), and a superficial gas velocity of approximately 0.6 m/s (2 ft/s). Total reactor pressure was 1.86 MPa gauge (270 psig). A seed bed was charged to the reactor and it was dried to 50 ppm water. It was pressurized to 0.69 MPa gauge (100 psig) of nitrogen and then 36 kg/hr (80 lbs/hr) of 10% TEAL in isopentane were to the reactor over two hours and allowed to circulate for 1 hour. The hexene-ethylene ratio was established at 0.028 and the temperature was adjusted. Atmer-163, a chemical marketed by ICI, was added as necessary to the reactor to control the buildup of electrostatic charge.

[0063] Catalyst was mixed with MMAO 3A as received at 7.38 wt % Al. Additional dilution was performed by adding isopentane to the mixture.

[0064] This liquid catalyst was introduced to the reactor with 6.4mm (1/4") OD stainless-steel injection tube of 4.6mm (0.18 inch) ID. For Examples 8 and 9, a 7.6cm (3-inch) length of tubing with a smaller ID was welded onto the end of the injection tube. The tube was located within a particle deflecting gas tube in Examples 8 and 9. For Examples 6, 7, and 8, the tip-cleaning gas tube extended 54 cm (23 inches) into the reactor at an elevation 6 inches above the distributor plate. For Example 9, the tip-cleaning gas tube extended 81 cm (32 inches) into the reactor at 2.4m (8 feet) above the plate. The injection tube was centered in the particle deflecting gas tube with tapered fins, and the tip of the injection tube extended 5 cm (2 inches) into the reactor beyond the end of the tip-cleaning gas tube. The ends of the injection tubes were tapered, to avoid accumulation of catalyst solution. The tip-cleaning gas tube was also tapered over the last 15 cm (6 inches). A stream of ethylene was fed through the annular space between the tip-cleaning gas tube and the catalyst injection tube. Example 8 demonstrated that stable particle size can be achieved with a combination of ethylene flowrate through the tip-cleaning gas tube, and with a high feed rate of nitrogen through the injection tube.

[0065] Table III illustrates of the present invention in polymerization processes that produce different polymers under differing reaction conditions.

TABLE II

	Ex. 6	Ex. 7	Ex. 8	Ex. 9
Ethylene flow kg/hr(lbs/hr)**	450 (1000)	900 (2000)	900 (2000)	230 (500)
Ethylene velocity m/s(ft/s)	20.4 (67)	40.5 (133)	39.9 (131)	83.8 (275)
Ethylene temperature(°C)	15	45	85	15
Shroud ID mm(inches)	21.5 (0.847)	21.5 (0.847)	21.5 (0.847)	9.53 (0.375)
Shroud OD mm(inches)	49.15 (1.935)	49.15 (1.935)	49.15 (1.935)	25.4 (1)
Shroud length cm(inches)	58 (23)	58 (23)	58 (23)	81 (32)
N ₂ flow kg/hr(lbs/hr)††	23 (50)	36 (80)	50 (110)	45 (100)
N ₂ velocity m/s(ft/s)	20.4(67)	32.9 (108)	100 (328)	94.8 (311)
Injector ID mm(inches)	4.6 (0.18)	4.6 (0.18)	3.0 (0.12)	2.98 (0.1175)
Injector OD mm(inches)	6.4(0.25)	6.4(0.25)	6.4(0.25)	6.4(0.25)
MMAO kg/hr (lbs/hr)	6.4 to 12.7 (14 to 28)	6.8 to 8.2 (15 to 18)	2.7 to 4.5 (6 to 10)	4.1 (9)
Catalyst kg/hr(lbs/hr)	0.9 to 1.8 (2 to 4)	0.2 to 0.5 (0.5 to 1)	0.5 to 0.8 (1 to 1.7)	0.5 (1)
Al:Zr (molar)	350	1000	350	350
Isopentane kg/hr(lbs/hr)	0	4.1 to 11 (9 to 24)	3.6 (8)	3.6 (8)
Temperature (°C)	80	80	75	75
C2 Partial Pressure MPa (psia)	1.28 to 1.38 (185 to 200)	1.17 to 1.24 (170 to 180)	1.17 to 1.24 (170 to 180)	1.17 to 1.24 (170 to 180)
Elevation above distributor m(ft)	0.15 (0.5)	0.15 (0.5)	0.15 (0.5)	2.4 (8)
Particle size mm(inches)	Increased rapidly to 1.98 (0.078)	Increased rapidly to 1.68 (0.066)	Stable at 0.81 (0.032)	Grew rapidly Steady state not reached

** Used as a carrier gas.

†† Used as a particle deflecting gas and as a tip cleaning gas.

TABLE III

	Ex. 10	Ex. 11	Ex. 12	Ex. 13	Ex. 14
Polymer	PE	PP	EPR	BR	Polyisoprene
Liquid Catalyst	Zr metallocene	Ti metallocene	V(AcAc) ₃	Nd Versatate	Nd neodecanate
Cocatalyst	MAO	MMAO	DEAC	TIBA	TEAL
Promoter	ETCA	DIBAH	MAO
Inert Particulate Material	Carbon black	Carbon black	Carbon black
Reactor Temperature °C	75	80	40	50	35
Reactor pressure MPa(psig)	2.07(300)	3.1(450)	2.76(400)	2.07(300)	2.07(300)
Gas Velocity m/s(ft/s)	0.61(2.0)	0.61(2.0)	0.52(1.7)	0.52(1.7)	0.52(1.7)
Space/Time/Yield kg/hr/m ³ (lb/hr/ft ³)	128(8)	128(8)	64(4)	64(4)	64(4)
Gas Composition					
Partial Pressure					
C ₂ H ₄	250	...	100
C ₃ H ₆	...	375	250
C ₄ H ₆
C ₃ H ₈	40	20
H ₂ /C ₃	0.01	...	0.025
H ₂ /C ₁	...	0.01
C ₁ /C ₃	2.5
Particle Deflecting Gas	Recycle Gas	Recycle Gas	Recycle Gas	Recycle Gas	Recycle Gas
Tip Cleaning Gas	Ethylene	Ethylene	Ethylene	Recycle Gas	Recycle Gas
Injection Means*	2	2	2	2	2

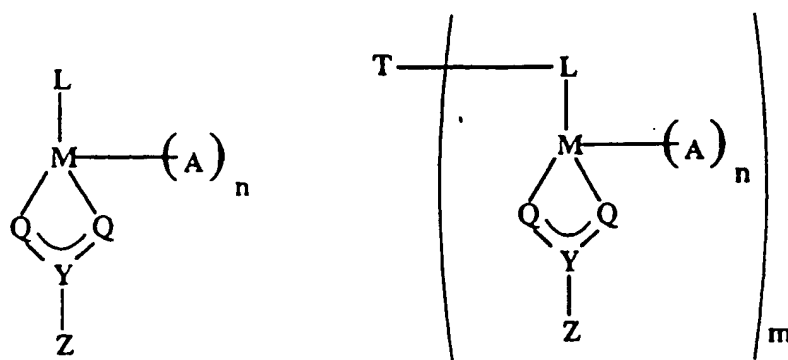
* 1 = Extended tapered air-cap spray nozzle (Fig. 2); 2 = Tapered injection tube (Fig. 3); 3 = Catalyst spray nozzle in purging shroud.

Claims

1. A process for producing a polymer in a gas fluidized bed reaction vessel having a polymerization zone containing

a bed of growing polymer particles under polymerization conditions which comprises:

- (i) continuously or intermittently introducing one or more monomers, optionally at least one diene, optionally at least one inert particulate material, and optionally one or more inert gases into the polymerization zone;
 - (ii) continuously or intermittently introducing into the reaction vessel an unsupported polymerization catalyst in liquid form without immediately contacting the bed of growing polymer particles so as to provide a particle lean zone;
 - (iii) continuously or intermittently withdrawing polymeric product from the reaction vessel; and
 - (iv) continuously withdrawing unreacted monomers and gases from said polymerization and optionally compressing and cooling said gases while maintaining the temperature within the polymerization zone below the dew point of at least one monomer present in the polymerization zone.
2. The process of Claim 1 wherein the droplets of the unsupported catalyst in liquid form are introduced without immediately contacting growing polymer particles of the bed so as to provide an average polymer particle size of 0.25 to 1.52 mm (0.01 to 0.06 inches).
3. The process of Claim 2 wherein the liquid catalyst is in a carrier gas surrounded by at least one particle-deflecting gas having a velocity sufficient to deflect resin particles of the bed away from the path of the liquid catalyst as it enters the fluidization zone.
4. The process of Claim 3 wherein the liquid catalyst is in a carrier gas surrounded by at least one particle-deflecting gas and at least one tip-cleaning gas, each gas selected from the group consisting of recycle gas, monomer gas, chain transfer gas, and inert gas.
5. The process of Claim 4 wherein the liquid catalyst in the carrier gas is introduced at a velocity of 15 to 120 m/s (50 to 400 ft/s); the particle deflecting gas is introduced at a velocity of 3 to 46 m/s (10 to 150 ft/s); and the tip-cleaning gas is introduced at a velocity of 15 to 76 m/s (50 to 250 ft/s).
6. The process of Claim 4 or 5 wherein the particle-deflecting gas and the tip-cleaning gas is 70 to 340 kPa gauge (10 to 50 psig) higher in pressure than the gas in the polymerization zone of the reaction vessel.
7. The process of any one of Claims 4 to 6 wherein the particle-deflecting gas is recycle gas and the tip-cleaning gas is monomer gas.
8. The process of any one of claims 4 to 7 wherein the liquid catalyst is introduced by a catalyst delivery system comprising (i) a particle-deflecting gas tube enclosing a catalyst injection tube or nozzle or (ii) a particle-deflecting gas tube enclosing a tip-cleaning gas tube which in turn encloses a catalyst injection tube or nozzle.
9. The process of any one of the preceding Claims wherein the liquid catalyst is selected from the group consisting of a metallocene catalyst, a rare earth catalyst, and mixtures thereof.
10. The process of Claim 9 wherein the metallocene catalyst composition comprises
 - (i) a catalyst precursor selected from the group consisting of



and mixtures thereof
wherein:

M is Zr or Hf;

L is a substituted or unsubstituted, π -bonded ligand;

each Q can be the same or different and is independently selected from the group consisting of -O-, -NR³-, -CR³₂- and -S-;

Y is either C or S;

Z is selected from the group consisting of -OR³, -NR³₂, -CR³₃, -SR³, -SiR³₃, -PR³₂ and -H, with the proviso that when Q is -NR³- then Z is selected from the group consisting of -OR³, -NR³₂, -SR³, -SiR³₃, -PR³₂ and -H;

\underline{n} is 1 or 2;

A is a univalent anionic group when \underline{n} is 2 or A is a divalent anionic group when \underline{n} is 1;

R³ can be the same or different and is independently a group containing carbon, silicon, nitrogen, oxygen, and/or phosphorus and one or more R³ groups may optionally be attached to the L substituent;

T is a bridging group selected from the group consisting of an alkylene or arylene group containing from 1 to 10 carbon atoms, germanium, silicon and alkyl phosphine; and \underline{m} is 1 to 7; and

(ii) a co-catalyst.

11. The process of Claim 9 or 10 wherein the rare earth catalyst is selected from the group consisting of a neodymium carboxylate, a neodymium alcoholate, a neodymium acetylacetonate, a neodymium halide, a neodymium allyl derivative, and mixtures thereof.
12. The process of any one of the preceding Claims wherein the polymer being produced is selected from the group consisting of (a) ethylene homopolymers, ethylene copolymers, and ethylene terpolymers employing one or more C₃-C₁₂ alpha olefins; (b) propylene homopolymers and propylene copolymers employing one or more C₄-C₁₂ alpha olefins; (c) polybutadiene; and (d) ethylene propylene rubbers and ethylene propylene diene rubbers.
13. The process of any one of the preceding Claims wherein the diene selected from the group consisting of 5-ethylidene-2-norbornene, 1,3-cyclopentadiene, 1,7-octadiene, 1,4-hexadiene, and mixtures thereof is employed.
14. The process of any one of the preceding Claims wherein the inert particulate material selected from the group consisting of carbon black, silica, clay, talc, and mixtures thereof is employed.

Patentansprüche

1. Verfahren zur Herstellung eines Polymers in einem Gas-Fließbett-Reaktionsgefäß mit einer Polymerisationszone, die ein Bett aus wachsenden Polymerteilchen enthält, unter Polymerisationsbedingungen, welches umfaßt:

(i) kontinuierliches oder absatzweises Einführen eines oder mehrerer Monomere, gegebenenfalls mindestens eines Diens, gegebenenfalls mindestens eines inerten teilchenförmigen Materials, und gegebenenfalls eines

oder mehrerer Inertgase in die Polymerisationszone;

(ii) kontinuierliches oder absatzweises Einführen eines nicht-geträgerten Polymerisationskatalysators in flüssiger Form in das Reaktionsgefäß ohne direktes Kontaktieren des Betts aus wachsenden Polymerteilchen, um eine Teilchen-arme Zone bereitzustellen;

(iii) kontinuierliche oder absatzweise Entnahme von Polymerprodukt aus dem Reaktionsgefäß; und

(iv) kontinuierlicher Abzug von nicht umgesetzten Monomeren und Gasen aus der Polymerisation und gegebenenfalls Verdichten und Abkühlen der Gase, während die Temperatur innerhalb der Polymerisationszone unterhalb des Taupunkts mindestens eines in der Polymerisationszone vorhandenen Monomers gehalten wird.

2. Verfahren nach Anspruch 1, worin die Tröpfchen des nicht-geträgerten Katalysators in flüssiger Form ohne direktes Kontaktieren der wachsenden Polymerteilchen des Betts eingeführt werden, um eine durchschnittliche Polymer-Teilchengröße von 0,25 bis 1,52 mm (0,01 bis 0,06 Zoll) bereitzustellen.

3. Verfahren nach Anspruch 2, worin sich der flüssige Katalysator in einem Trägergas befindet, das von mindestens einem Teilchen-ablenkenden Gas umgeben ist, welches eine Geschwindigkeit hat, die ausreicht, um Harzteilchen aus dem Bett vom Weg des flüssigen Katalysators bei dessen Eintritt in die Fluidisationszone abzulenken.

4. Verfahren nach Anspruch 3, worin sich der flüssige Katalysator in einem Trägergas befindet, das von mindestens einem Teilchen-ablenkenden Gas und mindestens einem Düsen spitzen-reinigenden Gas umgeben ist, wobei jedes Gas ausgewählt ist aus der Gruppe bestehend aus Kreisgas, Monomergas, Kettenübertragungsgas und Inertgas.

5. Verfahren nach Anspruch 4, worin der flüssige Katalysator in dem Trägergas mit einer Geschwindigkeit von 15 bis 120 m/s (50 bis 400 Fuß/s) eingeführt wird; das Teilchen-ablenkende Gas mit einer Geschwindigkeit von 3 bis 46 m/s (10 bis 150 Fuß/s) eingeführt wird; und das Düsen spitzen-reinigende Gas mit einer Geschwindigkeit von 15 bis 76 m/s (50 bis 250 Fuß/s) eingeführt wird.

6. Verfahren nach Anspruch 4 oder 5, worin das Teilchen-ablenkende Gas und das Düsen spitzen-reinigende Gas einen um 70 bis 340 kPa (10 bis 50 psig) höheren Überdruck hat als das Gas in der Polymerisationszone des Reaktionsgefäßes.

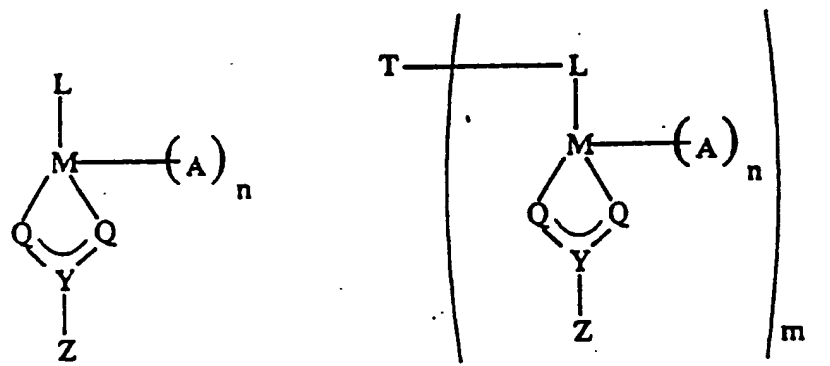
7. Verfahren nach irgendeinem der Ansprüche 4 bis 6, worin das Teilchen-ablenkende Gas Kreisgas ist und das Düsen spitzen-reinigende Gas Monomergas ist.

8. Verfahren nach irgendeinem der Ansprüche 4 bis 7, worin der flüssige Katalysator mit Hilfe eines Katalysator-Zufuhrsystems eingeführt wird, welches umfaßt (i) ein Rohr für Teilchen-ablenkendes Gas, welches eine Katalysator-Einspritzröhre oder -düse umschließt, oder (ii) ein Rohr für Teilchen-ablenkendes Gas, welches eine Röhre für Düsen spitzen-reinigendes Gas umschließt, die wiederum eine Katalysator-Einspritzröhre oder -düse umschließt.

9. Verfahren nach irgendeinem der vorhergehenden Ansprüche, worin der flüssige Katalysator ausgewählt ist aus der Gruppe bestehend aus einem Metallocen-Katalysator, einem Seltenerd-Katalysator und Mischungen davon.

10. Verfahren nach Anspruch 9, worin die Metallocen-Katalysatorzusammensetzung umfaßt

(i) eine Katalysator-Vorstufe, welche ausgewählt ist aus der Gruppe bestehend aus



und Mischungen davon, worin:

M Zr oder Hf bedeutet;

L ein substituierter oder unsubstituierter π -gebundener Ligand ist; jedes Q gleich oder verschieden sein kann und unabhängig ausgewählt ist aus der Gruppe bestehend aus -O-, -NR³-, -CR³₂- und -S-;

Y entweder C oder S bedeutet;

Z ausgewählt ist aus der Gruppe bestehend aus -OR³, -NR³₂, -CR³₃, -SR³, -SiR³₃, -PR³₂ und -H, mit der Maßgabe, daß wenn Q -NR³- ist, dann Z ausgewählt ist aus der Gruppe bestehend aus -OR³, -NR³₂, -SR³, -SiR³₃, -PR³₂ und -H;

n 1 oder 2 beträgt;

A eine einwertige anionische Gruppe bedeutet, wenn n 2 ist, oder A eine zweiwertige anionische Gruppe bedeutet, wenn n 1 ist;

R³ gleich oder verschieden sein kann und unabhängig eine Kohlenstoff, Silicium, Stickstoff, Sauerstoff und/oder Phosphor enthaltende Gruppe bedeutet und eine oder mehrere Gruppen R³ gegebenenfalls an den L-Substituenten gebunden sein können;

T eine Brückengruppe bedeutet, die ausgewählt ist aus der Gruppe bestehend aus einer 1 bis 10 Kohlenstoffatome enthaltenden Alkyl- oder Arylgruppe, Germanium, Silicon und Alkylphosphin; und m 1 bis 7 beträgt; und

(ii) einen Cokatalysator.

11. Verfahren nach Anspruch 9 oder 10, worin der Seltenerd-Katalysator ausgewählt ist aus der Gruppe bestehend aus einem Neodymcarboxylat, einem Neodymalkoholat, einem Neodymacetylacetonat, einem Neodymhalogenid, einem Neodymallylderivat und Mischungen davon.

12. Verfahren nach irgendeinem der vorhergehenden Ansprüche, worin das hergestellte Polymer ausgewählt ist aus der Gruppe bestehend aus (a) Ethylen-Homopolymeren, Ethylen-Copolymeren und Ethylen-Terpolymeren unter Verwendung eines oder mehrerer C₃-C₁₂- α -Olefine; (b) Propylen-Homopolymeren und Propylen-Copolymeren unter Verwendung eines oder mehrerer C₄-C₁₂- α -Olefine; (c) Polybutadien; und (d) Ethylen-Propylen-Kautschuken und Ethylen-Propylen-Dien-Kautschuken.

13. Verfahren nach irgendeinem der vorhergehenden Ansprüche, worin das aus der Gruppe bestehend aus 5-Ethyliden-2-norbornen, 1,3-Cyclopentadien, 1,7-Octadien, 1,4-Hexadien und Mischungen davon ausgewählte Dien verwendet wird.

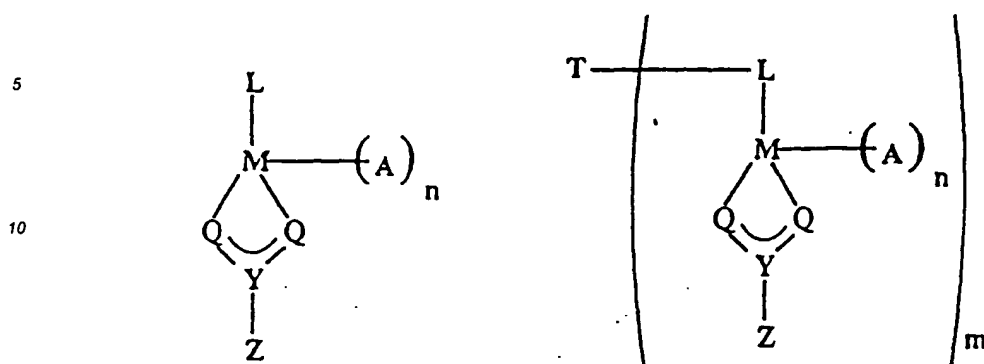
14. Verfahren nach irgendeinem der vorhergehenden Ansprüche, worin das aus der Gruppe bestehend aus Ruß, Kieselsäure, Tonerde, Talkum und Mischungen davon ausgewählte inerte teilchenförmige Material verwendet wird.

Revendications

1. Procédé pour la production d'un polymère dans un récipient de réaction à lit fluidisé en phase gazeuse, comprenant une zone de polymérisation contenant un lit de particules de polymère en croissance, dans des conditions de

polymérisation, qui comprend les étapes consistant :

- (i) à introduire de manière continue ou intermittente un ou plusieurs monomères, facultativement au moins un diène, facultativement au moins une matière inerte en particules et facultativement un ou plusieurs gaz inertes dans la zone de polymérisation ;
 - (ii) à introduire de manière continue ou intermittente dans le récipient de réaction un catalyseur de polymérisation non fixé sur un support, sous forme liquide, sans mettre immédiatement en contact le lit de particules de polymère en croissance de manière à produire une zone pauvre en particules ;
 - (iii) à décharger de manière continue ou intermittente le produit polymère du récipient de réaction ; et
 - (iv) à décharger de manière continue les monomères et gaz n'ayant pas réagi de ladite zone de polymérisation et, facultativement, à comprimer et refroidir lesdits gaz tout en maintenant la température à l'intérieur de la zone de polymérisation à une valeur inférieure au point de rosée d'au moins un monomère présent dans la zone de polymérisation.
2. Procédé suivant la revendication 1, dans lequel les gouttelettes du catalyseur non fixé sur un support, sous forme liquide, sont introduites sans mettre immédiatement en contact les particules de polymère en croissance du lit de manière à parvenir à un diamètre moyen de particules de polymère de 0,25 à 1,52 mm (0,01 à 0,06 inch).
 3. Procédé suivant la revendication 2, dans lequel le catalyseur liquide est présent dans un gaz servant de véhicule entouré par au moins un gaz de déflexion de particules ayant une vitesse suffisante pour dévier les particules de résine du lit hors du trajet du catalyseur liquide lors de son entrée dans la zone de fluidisation.
 4. Procédé suivant la revendication 3, dans lequel le catalyseur liquide est présent dans un gaz servant de véhicule entouré par au moins un gaz de déflexion de particules et au moins un gaz de nettoyage d'embout, chaque gaz étant choisi dans le groupe consistant en un gaz de recyclage, un monomère gazeux, un gaz de transfert de chaîne et un gaz inerte.
 5. Procédé suivant la revendication 4, dans lequel le catalyseur liquide présent dans le véhicule gazeux est introduit à une vitesse de 15 à 120 m/s (50 à 400 ft/s) ; le gaz de déflexion de particules est introduit à une vitesse de 3 à 46 m/s (10 à 150 ft/s) ; et le gaz de nettoyage d'embout est introduit à une vitesse de 15 à 76 m/s (50 à 250 ft/s).
 6. Procédé suivant la revendication 4 ou 5, dans lequel le gaz de déflexion de particules et le gaz de nettoyage d'embout ont une pression supérieure de 70 à 340 kPa au manomètre (10 à 50 psig) à la pression du gaz dans la zone de polymérisation du récipient de réaction.
 7. Procédé suivant l'une quelconque des revendications 4 à 6, dans lequel le gaz de déflexion de particules consiste en gaz de recyclage et le gaz de nettoyage d'embout consiste en monomère gazeux.
 8. Procédé suivant l'une quelconque des revendications 4 à 7, dans lequel le catalyseur liquide est introduit par un système de distribution de catalyseur comprenant (i) un tube de gaz de déflexion de particules entourant un tube ou une buse d'injection de catalyseur ou (ii) un tube de gaz de déflexion de particules entourant un tube de gaz de nettoyage d'embout qui entoure lui-même un tube ou une buse d'injection de catalyseur.
 9. Procédé suivant l'une quelconque des revendications précédentes, dans lequel le catalyseur liquide est choisi dans le groupe consistant en un catalyseur à base de métallocène, un catalyseur à base de terre rare et leurs mélanges.
 10. Procédé suivant la revendication 9, dans lequel la composition de catalyseur à base de métallocène comprend
(i) un précurseur de catalyseur choisi dans le groupe consistant en



et leurs mélanges
formules dans lesquelles :

M représente Zr ou Hf ;

L représente un ligand à liaison n, substitué ou non substitué ;

les groupes Q peuvent être identiques ou différents et sont choisis chacun, indépendamment, dans le groupe consistant en -O-, -NR³-, -CR³₂- et -S- ;

Y représente C ou S ;

Z est choisi dans le groupe consistant en -OR³, -NR³₂, -CR³₃, -SR³, -SiR³₃, -PR³₂ et -H, sous réserve que, lorsque Q représente un groupe -NR³-, alors Z soit choisi dans le groupe consistant en -OR³, -NR³₂, -SR³, -SiR³₃, -Pr³₂ et -H ;

n est égal à 1 ou 2 ;

A représente un groupe anionique univalent lorsque n est égal à 2 ou bien A représente un groupe anionique divalent lorsque n est égal à 1 ;

les groupes R³ peuvent être identiques ou différents et représentent chacun, indépendamment, un groupe contenant du carbone, du silicium, de l'azote, de l'oxygène et/ou du phosphore, et un ou plusieurs groupes R³ peuvent être facultativement fixés au substituant L ;

T représente un groupe de pontage choisi dans le groupe consistant en un groupe alkylène ou arylène contenant 1 à 10 atomes de carbone, le germanium, le silicium et une alkylphosphine ; et m a une valeur de 1 à 7 ; et

(ii) un cocatalyseur.

11. Procédé suivant la revendication 9 ou 10, dans lequel le catalyseur à base de terre rare est choisi dans le groupe consistant en un carboxylate de néodyme, un alcoolate de néodyme, un acétylacétonate de néodyme, un halogénure de néodyme, un dérivé allylique de néodyme et leurs mélanges.

12. Procédé suivant l'une quelconque des revendications précédentes, dans lequel le polymère produit est choisi dans le groupe consistant en (a) des homopolymères d'éthylène, des copolymères d'éthylène et des terpolymères d'éthylène utilisant une ou plusieurs alpha-oléfines en C₃ à C₁₂ ; (b) des homopolymères de propylène et copolymères de propylène utilisant une ou plusieurs alpha-oléfines en C₄ à C₁₂ ; (c) le polybutadiène ; et (d) des caoutchoucs éthylène-propylène et des caoutchoucs éthylène-propylène-diène.

13. Procédé suivant l'une quelconque des revendications précédentes, dans lequel on utilise le diène choisi dans le groupe consistant en le 5-éthylidène-2-norbornène, le 1,3-cyclopentadiène, le 1,7-octadiène, le 1,4-hexadiène et leurs mélanges.

14. Procédé suivant l'une quelconque des revendications précédentes, dans lequel on utilise la matière inerte en particules choisie dans le groupe consistant en le noir de carbone, la silice, une argile, le talc et leurs mélanges.

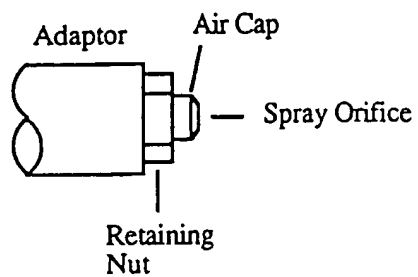


Figure 1. Profile of commercially available air cap mounted in custom adapter.

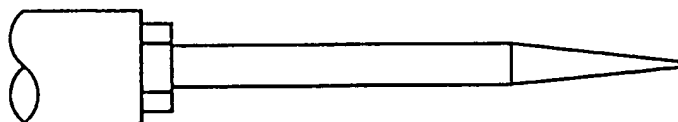


Figure 2. Extended tapered "air-cap" for catalyst spray nozzle.

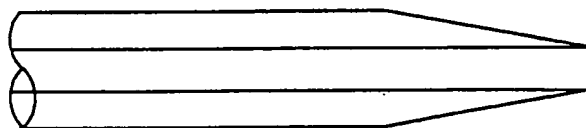


Figure 3. Tapered injection tube tip.

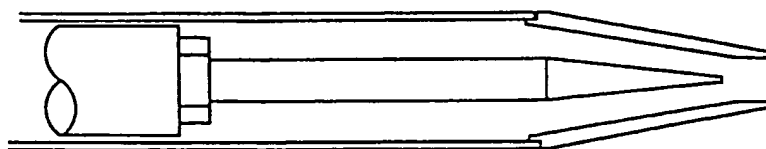


Figure 4. Catalyst spray nozzle enclosed in purging shroud.